## In the claims:

- 28. (Amended Twice) A method for enhancing the formation and development of dendrites and synapses in hippocampal neurons, comprising contacting said neurons with a morphogen selected from: an OP-1 polypeptide, a BMP-2 polypeptide, a BMP-5 polypeptide, a BMP-6 polypeptide, or a 60A polypeptide, wherein said morphogen has a conserved C-terminal seven-cysteine skeleton at least about 60% identical to residues 330-431 of human OP-1 (SEQ ID NO: 2), and wherein said morphogen induces dendrite outgrowth in said hippocampal neuron.
  - 29. (Reiterated) The method of claim 28, wherein said morphogen comprises residues 30-292 of SEQ ID NO: 2.
  - 30. (Reiterated) The method of claim 28, wherein said morphogen comprises residues 330-431 of SEQ ID NO: 2.
  - 31. (Reiterated) The method of claim 28, wherein said morphogen comprises residues 48-292 of SEQ ID NO: 2.
  - 32. (Reiterated) The method of claim 28, wherein said morphogen comprises the amino acid sequence of SEQ ID NO: 2.

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- 46. (Amended) The method of claim 28, wherein said morphogen comprises residues 293-329 of SEQ ID NO: 2.
- 47. (Amended) The method of claim 28, wherein said morphogen comprises residues 293-431 of SEQ ID NO: 2.
- 48. (Reiterated) The method of claim 28, wherein said morphogen comprises residues 30-431 of SEQ ID NO: 2.

Please add the following new claims:

- 52. (New) A method for enhancing the formation and development of dendrites and synapses in hippocampal neurons, comprising contacting said neurons with a morphogen selected from: an OP-1 polypeptide, a BMP-2 polypeptide, a BMP-5 polypeptide, a BMP-6 polypeptide, or a 60A polypeptide, wherein said morphogen has a conserved C-terminal seven-cysteine skeleton at least about 70% homologous to residues 330-431 of human OP-1 (SEQ ID NO: 2), and wherein said morphogen induces dendrite outgrowth in said hippocampal neuron.
- 53. (New) The method of claim 52, wherein said morphogen comprises residues 30-292 of SEQ ID NO: 2.
- 54. (New) The method of claim 52, wherein said morphogen comprises residues 330-431 of SEQ ID NO: 2.
- 55. (New) The method of claim 52, wherein said morphogen comprises residues 48-292 of SEQ ID NO: 2.
- 56. (New) The method of claim 52, wherein said morphogen comprises the amino acid sequence of SEQ ID NO: 2.
- 57. (New) The method of claim 52, wherein said morphogen comprises residues 293-329 of SEQ ID NO: 2.
- 58. (New) The method of claim 52, wherein said morphogen comprises residues 293-431 of SEQ ID NO: 2.
- 59. (New) The method of claim 52, wherein said morphogen comprises residues 30-431 of SEQ ID NO: 2.

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The claims presented above incorporate changes as indicated by the marked-up versions below.

- 28. (Amended Twice) A method for enhancing the formation and development of dendrites and synapses in hippocampal neurons eells, comprising contacting said neurons eells with a morphogen selected from: an OP-1 polypeptide, a BMP-2 polypeptide, a BMP-5 polypeptide, a BMP-6 polypeptide, and or a 60A polypeptide, wherein said morphogen has a conserved C-terminal seven-cysteine skeleton at least about 60% identical to residues 330-431 of human OP-1 (SEQ ID NO: 2), and wherein said morphogen induces dendrite outgrowth in said hippocampal neuron.
- 46. (Amended) The method of claim 28, wherein said morphogen comprises residues <del>292-</del> <del>293-329</del> of SEQ ID NO: 2.
- 47. (Amended) The method of claim 28, wherein said morphogen comprises residues 292-431 of SEQ ID NO: 2.